AMENDMENTS TO THE CLAIMS

1. (Currently amended) A method for producing a synchronized population of

conifer somatic embryos, the method comprising the step of cultivating pre-cotyledonary conifer

embryogenic cells in, or on, a synchronization medium that comprises an absorbent composition

and at least one synchronization agent selected from the group consisting of abscisic acid and a

gibberellin, wherein the absorbent composition and the at least one synchronization agent are

present at a concentration effective to produce a synchronized population of pre-cotyledonary

conifer somatic embryos, wherein the pre-cotyledonary conifer embryogenic cells are cultivated

in the synchronization medium prior to cultivation in a development medium.

2. (Original) The method of Claim 1 wherein the absorbent composition is selected

from the group consisting of activated charcoal, soluble poly(vinyl pyrrolidone), insoluble

poly(vinyl pyrrolidone), activated alumina, and silica gel.

3. (Original) The method of Claim 2 wherein the absorbent composition is activated

charcoal.

4. (Original) The method of Claim 1 wherein the concentration of the absorbent

composition in the synchronization medium is from about 0.5 g/L to about 50 g/L.

5. (Original) The method of Claim 1 wherein the absorbent composition is activated

charcoal, and the activated charcoal is present in the synchronization medium at a concentration

in the range of from about 0.1 g/L to about 5 g/L.

(Original) The method of Claim 1 wherein the absorbent composition is activated

charcoal, and the activated charcoal is present in the synchronization medium at a concentration

in the range of from about 0.5 g/L to about 1 g/L.

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6.

(Original) The method of Claim 1, wherein abscisic acid is used as a 7. synchronization agent.

(Original) The method of Claim 1, wherein a gibberellin is used as a 8.

synchronization agent.

(Original) The method of Claim 1, wherein abscisic acid and at least one 9.

gibberellin are used as synchronization agents.

(Original) The method of Claim 1, wherein a gibberellin is present in the 10.

synchronization medium at a concentration of from about 0.5 mg/L to about 500 mg/L.

(Original) The method of Claim 1, wherein a gibberellin is present in the 11.

synchronization medium at a concentration of from about 1.0 mg/L to about 100 mg/L.

(Original) The method of Claim 1, wherein abscisic acid is present in the 12.

synchronization medium at a concentration of from about 1.0 mg/L to about 500 mg/L.

(Original) The method of Claim 1, wherein abscisic acid is present in the 13.

synchronization medium at a concentration of from about 0.5 mg/L to about 20 mg/L.

(Original) The method of Claim 1, wherein the conifer embryogenic cells are 14.

cultured in, or on, the synchronization medium for a period of from about 0.5 weeks to about

5 weeks.

(Original) The method of Claim 1, wherein the conifer embryogenic cells are 15.

cultured in, or on, the synchronization medium for a period of from about 1 week to about

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3 weeks.

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(Original) The method of Claim 1, wherein the conifer embryogenic cells are 16. cultured in, or on, the synchronization medium for a period of from about 1 week to about

2 weeks.

(Original) The method of Claim 1, wherein the osmolality of the synchronization 17.

medium is from about 90 mM/Kg to about 300 mM/Kg.

(Original) The method of Claim 1, wherein the pH of the synchronization 18.

medium is from about 5 to about 6.

(Original) The method of Claim 1, wherein Loblolly pine somatic embryos are 19.

produced from Loblolly pine embryogenic cells.

(Original) The method of Claim 1, wherein at least 50% of the embryos in the 20.

synchronized population of conifer somatic embryos are at the same developmental stage.

(Original) The method of Claim 1, wherein at least 75% of the embryos in the 21.

synchronized population of conifer somatic embryos are at the same developmental stage.

(Currently amended) The method of Claim 1, wherein the synchronized 22.

population of pre-cotyledonary conifer somatic embryos are transferred to [[a]] the development

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medium for synchronized cotyledonary embryo development.

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